On the Function of Ceria Supported Rhodium Catalysts for Methanation of Carbon Dioxide

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Department of Chemistry and Chemical Engineering
Chalmers University of Technology
Gothenburg, Sweden 2020
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Abstract

Technologies for energy harvesting of sustainable sources such as solar and wind lack an inherent energy buffer. As such, storing excess energy in chemical bonds, for example that of hydrogen (H₂), is a desirable complementary concept. While hydrogen gas is easily cycled and applied in electrical systems, it brings high costs for long term storage and transportation. The technical solutions introduce complications in terms of limited equipment lifetime and when applied in combustion systems also safety concerns. An alternative for these applications is to further convert the hydrogen into hydrocarbons such as alkanes and alcohols. Through CO₂ hydrogenation, methane can be produced and used in current infrastructure solutions due to it being the major component also in natural gas. While typically produced using nickel based catalysts, other metals such as ruthenium and rhodium have shown promise, of which Rh/CeO₂ is one such catalyst. While kinetic studies exist of said catalyst, the reaction mechanism is unknown, which hampers further development.

This work aims towards clarifying the catalyst structure-function relationship as well as the important surface reactions. The ambition is to facilitate the start of fundamental research approach, which can later be developed to achieve a more complete understanding, allowing for tuning the important properties of said catalyst.

In this work, Rh/CeO₂ catalysts were prepared by incipient wetness impregnation and studied in situ during CO₂ hydrogenation in regards to its structural behaviour, by the use of high-energy X-ray diffraction and ambient pressure X-ray photoelectron spectroscopy, as well as the surface reactions using diffuse reflectance infrared spectroscopy. As signals from processes and adsorbates that truly participate in the reaction may be hidden by spectator signals, these studies were performed transiently as to enhance the response of active species while decreasing that of spectators.

It was revealed that CeO₂ is active during reaction conditions, possibly partaking in a cycle of formation and healing of oxygen vacancies during the reaction cycle as evidenced by the cycling of Ce⁴⁺ and Ce³⁺. Furthermore, while predominantly reduced, a RhOₓ phase was observed, showing the strong metal support interaction of Rh and CeO₂. Regarding surface reactions, several carbonyl species (b-CO, h-CO, m-CO) were shown to be active on the catalyst, as well as some carbonates (b-CO₃, p-CO₃) and formate (b-HCOO) species.

While the complete pathway need more experimental data to be determined, the activity of the carbonyl species suggests that the reaction follows a carbon monoxide based pathway such as the carbide pathway.

Keywords: Sabatier reaction; CO₂ reduction; Catalytic methanation; Hydrogenation; In situ spectroscopy; HE-XRD; AP-XPS; DRIFTS
List of Publications

This thesis is based on the following appended papers:

I. Structure-function relationship for CO\textsubscript{2} methanation over ceria supported Rh and Ni catalysts at atmospheric pressure conditions


*Catalysis Science & Technology, 9 (2019), 1644-1653*

II. CO\textsubscript{2} Methanation over Rh/CeO\textsubscript{2} Studied with Infrared Modulation Excitation Spectroscopy and Phase Sensitive Detection

F. Hemmingsson, A. Schaefer, M. Skoglundh and P.-A Carlsson

*Submitted manuscript (2020)*
My Contributions to the Publications

Paper I
I set-up and co-performed the HE-XRD experiments and co-authored the paper.

Paper II
I designed and performed all experiments and analyses, excluding XRF, and authored the manuscript.
Publications not included in the thesis:

**Structure–function relationship during CO$_2$ methanation over Rh/Al$_2$O$_3$ and Rh/SiO$_2$ catalysts under atmospheric pressure conditions**


*Catalysis Science & Technology, 8 (2018), 2686-2696*

**Methane Adsorption and Methanol Desorption of Copper Modified Boron Silicate**


*RSC Advances, 8 (2018), 3639-36374*
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The search for sustainable energy sources and systems has amassed a significant interest among societal actors, engineers and researchers. While the focus often, and perhaps foremost, has been on the availability of energy sources, an important shift towards questions regarding energy storage and fuels that are not sourced from fossil reserves has been witnessed during the latest decades [1, 2]. Though energy harvesting by wind and solar power shows great promise as “green” energy solutions [3], their dependence on external factors, e.g., weather, season and grid load, make them lack the necessary on-demand energy supply capability [4]. This currently limits their possibility of constituting a larger portion of countries’ energy system if not an energy buffer can be introduced. One such buffer could be the formation of gaseous hydrogen (H$_2$) by for instance water splitting (eq. (1.1)). Thereby the obtained energy is chemically stored in the hydrogen bond. This concept could be set up as a closed system by recycling the constituent molecules and is thus characterised by a zero net consumption/production of H$_2$O/H$_2$ [5].

\[
2\text{H}_2\text{O} \rightleftharpoons 2\text{H}_2 + \text{O}_2 \quad (1.1)
\]

Indeed, the concept of using H$_2$ as a basal energy vector has by itself gained much attention and the concept has been coined the hydrogen economy[6]. Even if H$_2$ is easily cycled and suitable applied in electrical systems, by the use of fuel cells [7], there are still a number of inherent problems associated with the broader use of H$_2$. For combustion systems, for example, a small H$_2$ concentration in the gas stream is tolerable but higher concentrations poses safety risks in terms of corrosion, i.e., hydrogen embrittlement, with consequential risk of leakage [9–11]. Furthermore, due to it being a gas at normal conditions, which is relevant for the majority applications, H$_2$ lacks the appropriate energy density and has a high storage cost in comparison to other fuels [12]. In order to circumvent these problems, it is possible to convert H$_2$ into alkanes and alcohols by hydrogenation of carbon oxides, i.e., carbon monoxide (CO) and carbon dioxide (CO$_2$) [2]. In contrast to H$_2$, long-chained alkanes and alcohols are liquids at normal conditions and can often be applied directly in existing systems such as combustion engines. However, while obtaining high energy densities and low storage costs, the processes for the production of these fuels incur an energy loss and require expensive high-

1: Furthermore, for certain applications an addition of hydrogen could prove beneficial, such as the case of utilisation of hydromethane (natural gas and hydrogen mixtures) [8].

2: Or with minor adjustments [13]. Furthermore, as these are base chemicals they are also of interest for applications other than fuel.

3: As an example, $\Delta H_f^o$(CH$_3$OH(g)) = −201.17 kJ mol$^{-1}$ [14].
In addition, some niche applications has arisen, such as the removal of CO\textsubscript{2} from air and recycling of H\textsubscript{2}O, as done at the international space station \cite{16}.

Figure 1.1: Schematic energy diagram for a reaction, showing the activation energy (\(E_a\)) for a non-catalysed reaction and a catalysed reaction. For the catalysed reaction, an apparent change in activation energy is shown as the dashed line whereas the full line shows the actual reaction with intermediary steps.

pressure chemical reactor systems. Further, while being suitable for long term storage and clearly usable for automotive-applications \cite{13,15}, the role of long-chained alkanes and alcohols for short-term energy cycling is questionable.

1.1 Catalysis for energy applications

The Sabatier reaction (equation (1.2)) was originally developed for removal of carbon oxides from hydrogen streams but has gained renewed interest in recent times thanks to the methane (CH\textsubscript{4}) product that is an attractive energy carrier\textsuperscript{4} \cite{12}.

\[
\text{CO}_2 + 4 \text{H}_2 \rightarrow \text{CH}_4 + 2 \text{H}_2\text{O}
\]  

(1.2)

Although CH\textsubscript{4} has lower energy density compared to long-chain alkanes and alcohols, its ease of production makes it suitable for intermediate storage times and short-term energy cycling \cite{17}. Furthermore, it can be utilised in current technologies developed for natural gas in regards to transport and usage, thus decreasing the implementation cost. Despite the fact that a chemical reaction might be energetically favoured, that alone may not warrant it being plausible. Take as an example diamonds. While De Beers slogan is a diamond is forever they are in fact unstable and degrade slowly forming graphite (\(\Delta G^\circ = -2.9 \text{ kJ mol}^{-1}\) \cite{18}). Fortunately, the degradation time amounts to billions of years, and as such you do not need to worry about your engagement ring slowly turning into a decorative pencil. This is because the degradation reaction is (strongly) kinetically limited, i.e., the reaction rate is sluggish, which is a characteristic that many industrially relevant reactions share. To speed up the kinetics one may, for example, heat up the sample because the rate of chemical reactions increase with increased temperature. The downside with heating is that it will not only increase the running cost of the reaction but might also open up for unwanted side reactions lowering the desired reaction selectivity. A fascinating approach to circumvent this problem is the usage of a catalyst. While the catalyst can not change the free energy of a reaction, i.e. a thermodynamically impossible reaction remains impossible, it can greatly increase its rate towards equilibrium by providing an alternative reaction pathway \cite[pp 1–4]{19}. Analogously, one can envision the catalytic process as taking a shortcut. We still move from point A to B (potential energy change is the same) but use less energy intense steps to do so. A more formal description is depicted in Figure 1.1. Here \(E_a\) is the activation energy of the reaction, the energy needed for the reaction to occur, and \(I_x\) are the intermediary steps of the reaction.
Like the shortcut, a catalyst is not consumed by its use. However, its shape and behaviour may change during the reaction cycle, but returns to the state of the initial step after completing the reaction cycle \([19, \text{ pp 1–4}]\).

### 1.2 Motivation of this work

The slow kinetics of catalysts for the Sabatier reaction makes high temperatures and pressures needed in order to gain industrially relevant yields \([20, 21]\). As such, the development of catalyst capable of operating close to ambient conditions would decrease the energy intensity of the process and open up for the methanation reaction to be run in less expensive industrial units and/or smaller remote apparatuses.
2.1 The Sabatier reaction

During the turn of the 20th century, Paul Sabatier together with Jean-Baptiste Senderens discovered what is now known as the Sabatier reaction, shown in equation (2.1), for which Sabatier was awarded the Nobel prize in chemistry in 1912 [22]. While originally utilised for purification of gaseous hydrogen from carbon oxides, it has now gained a renewed interest due to applications such as fuel production, energy storage and oxygen recycling [12].

\[
\text{CO}_2(g) + 4 \text{H}_2(g) \rightarrow \text{CH}_4(g) + 2 \text{H}_2\text{O}(g) \quad (2.1)
\]

\[
\Delta H_{298K} = -165 \text{ kJ mol}^{-1}
\]

The reaction is strongly exothermic and the drop in entropy limits the maximum conversion of the Sabatier reaction at atmospheric pressure, unless carried out at low temperatures. As such, industrially elevated temperature and pressure is utilised in order to achieve fast kinetics and to avoid the limitation to the maximum conversion [20, 21]. In regards to energy usage and simplicity of equipment, a low temperature and pressure solution might be more suitable. For heterogeneous catalysts operating at atmospheric conditions, initial activity at temperatures as low as 200 °C and operable temperatures (high conversion) as low as 300 °C are not uncommon for methanation catalysts [23, 24].

Ever since the days of Sabatier, methanation catalyst are predominantly composed of nickel, most often deposited on high surface metal oxide supports [12, 21, 25]. While metals such as rhodium and ruthenium can surpass nickel in regards to activity, the high activity and selectivity together with much lower price makes nickel based catalysts so far more favourable [21, 24]. However, some aspects limit the nickel catalysts such as deactivation by oxidation and formation of volatile carbonyls at low temperatures (< 300 °C), and carbon deposition [12, 20, 21, 26, 27]. For these reasons, noble metals are still of interest in the pursuit of decreased reaction temperature for CO₂ methanation.
Proposed reaction mechanisms

While a black box approach of catalyst screening allows for the discovery of new well performing catalysts using also some chemical intuition, it can seldom provide the insight needed to finely tune the properties of the catalyst. In order to accomplish this, knowledge of the reaction mechanism is needed in order to, for example, determine the rate determining step of the reaction and the cause behind its limitation. For CO₂ hydrogenation, as well as CO₂ hydrogenation in general, the reaction pathway is commonly following one of three different routes: the formate, carbon monoxide and carbonate pathway. Among these, it is the carbon monoxide and formate routes that have been observed experimentally, see equations (2.2)-(2.7) [12, 25]. For the traditional Ni catalysts, e.g., Ni/Sl₂O₃, Ni/Al₂O₃ and the like, which have enjoyed an extensive amount of studies, the carbide pathway in which the CO dissociates and forms elementary carbon (C) that is further hydrogenated into CH₄ is the most commonly attributed pathway [12, 21, 25]. However, while proceeding by a carbon monoxide based pathway, the intermediary steps of the nickel based catalysts have been shown to be dependent on the support. As an example, an IR study of a Ni/ZrO₂ catalyst where the ZrO₂ is an active support, intermediary bicarbonates and formates were observed, see equations (2.14)-(2.16) [25]. As such, while studying the reaction mechanism one may encounter an interplay between pathways as well as red herrings as the observation of intermediates most commonly associated to certain pathways might lead to the conclusion that the reaction follows that particular pathway.

2.2 The importance of transient in situ studies to reveal reaction mechanisms

While both ex situ and in situ studies in regards to the surface reactions are often conducted with ease by the use of methods such as infrared spectroscopy (see Section 3.4), the behaviour of the catalyst material during the reaction is not as easily studied, often leaving it as an ex situ comparison of the fresh and spent material. Crucial information such as the restructuring of the catalyst during the reaction therefore go unnoticed. Furthermore, not only active species but also spectators, a species that is formed at reaction
conditions but does not participate in the (main) reaction [29], may be observed and falsely attributed as a true reaction intermediate. Additional factors must therefore be included as to separate the contributions of active and spectators species, for which transient studies are common. This is further discussed in some depth in Section 3.5.

2.3 Objectives

This thesis aims at providing fundamental understanding of the behaviour of the Rh/CeO\(_2\) methanation catalyst under operation conditions and a first insight into the mechanistic pathway of the CO\(_2\) methanation reaction over this type of catalyst.
In this work, the \textit{in situ} behaviour of the catalyst material and the surface reactions over the catalyst for the considered chemical reaction were studied. To accomplish this, the changes in crystal structure and oxidation state was studied using synchrotron X-ray sources (Section 3.2 and 3.3) and the adsorbates of the reaction was studied by infrared spectroscopy (Section 3.4). As both active, those that participate in the reaction, and spectators, those that do not participate in the reaction, are observed in the resulting data, these experiments were conducted transiently using a modulation excitation approach (Section 3.5) in order to remove the contribution of spectators from the data set.

### 3.1 Catalyst preparation

One of the principle design rules for a heterogeneous (solid) catalyst to be efficient is that it should expose a high specific surface area (m$^2$/g) hosting the active sites or, as in the case of supported catalysts, carrying and stabilising the catalytically active particles\cite{19,29}. There are numerous ways of making catalysts, ranging from top-down approaches such as milling the active material into powder to bottom-up approaches with, for example, controlled and well defined nanoparticle synthesis and deposition\cite{31–34}. Another common route that is simple, yet effective, is impregnation of a support material with a solution of a precursor material, which gives small and well dispersed nanoparticles on that support\cite{32}. The impregnation can be made in different ways. The simplest form of wet impregnation uses an excess of precursor solution and the remaining liquid is evaporated by applying heat. The procedure may result in large particles and clogging of the pores of the support material, which decreases the material utilization and influences the catalytic properties of the catalyst. An alternative route is the dry impregnation, also called incipient wetness impregnation or capillary impregnation\cite{32}. The support material is impregnated with a liquid volume just enough to fill the pores. The sample is then frozen and freeze-dried as to remove the solvent by sublimation. This way, the problems of wet impregnation are avoided as the precursor is immobilised and largely not affected by drag forces by the solvent during removal\cite{32}.

![Catalyst preparation diagram](image-url)
In this thesis, the ceria supported rhodium (Rh/CeO$_2$) catalysts were prepared by incipient wetness impregnation, using Rh(NO$_3$)$_3$ (Paper I) or Rh dissolved in nitric acid (Paper II) as Rh precursor.

### 3.2 X-ray diffraction

A well known optical phenomenon is the bending of light when it passes through slits of size similar to or less than its wavelength. The concept is referred to as diffraction and can be used to determine the crystal structure of a material [35], which can be thought of as planes of atoms forming an infinite number of repeating slits. As typical inter-atomic distances are in the Ångström scale, X-rays provide the suitable wavelength to perform the experiment. The mathematical model for the description of the diffraction angle ($\theta$) in regards to the inter-planar distance ($d$) and wavelength ($\lambda$) was first presented by Bragg in 1913 [36]. The corresponding equation is referred to as the Bragg’s law and can be seen in equation (3.1) and a schematic representation of the process can be seen in Figure 3.2 [35].

\[
2d \sin(\theta) = n\lambda
\]  

Upon making a diffraction experiment, the obtained data is typically presented as a diffractogram, i.e., the measured intensity of the diffracted light against its corresponding diffraction angle as $2\theta$. As the angle of diffraction in equation (3.1) is wavelength dependent, one cannot directly compare diffractograms produced by different X-ray sources. In order to circumvent this problem, one may move from the R-space expression to the q-space, expressing it as a relation between the diffraction angle and wavelength, see equation (3.2).

\[
q = \frac{4\pi}{\lambda} \sin(\theta) = \frac{2\pi}{d}
\]  

The diffraction caused by a perfect single crystal, which has a defined orientation, results in discrete and clear diffraction spots [35]. For powder samples, ring patterns are instead produced due to the random orientation of the crystals in the analyzed powder [35]. Nevertheless, the diffraction angle is still the same.

In this study, the catalyst was studied by high energy X-ray diffraction (HE-XRD) with a high photon flux by the use of synchrotron light. The high energy and photon flux allowed for a detailed transient in situ study, which is prohibited by conventional lab X-ray sources.
3.3 X-ray photoelectron spectroscopy

The discovery of the photoelectric effect [37], the emission of (most commonly) an electron following the absorption of electromagnetic radiation, have lead to numerous different applications out of which X-ray photoelectron spectroscopy is one which is crucial in the field of catalysis research [19]. By measuring the kinetic energy of the ejected photoelectron one can find the original binding energy of said electron by equation (3.3), given that the energy of the X-ray photon (hv) and work function of the instrument (ϕ) is known [19]. This process is illustrated in Figure 3.3.

\[ E_{\text{binding}} = h\nu - E_{\text{kinetic}} + \phi \] (3.3)

This binding energy can not only reveal the kind of atoms present in the sample, but effects such as shifts of its peak in the spectrum may also provide an insight into its oxidation state.

Typically, XPS is limited to pressures ≤ 10⁻⁶ mbar due limitations to the path length of the emitted photoelectron by inelastic scattering by the surrounding gas phase [38]. As such, in situ studies by XPS have been severely limited in regards to catalysis studies. In more recent years, ambient pressure XPS (AP-XPS) in which the gas pressure of the sample cell is close to atmospheric condition have become possible thanks to sophisticated instrumental setup and high photon flux, supplied by synchrotron sources [38–40].

In this study, the oxidation state of the catalyst as well as the formation of some adsorbates formed was studied during CO₂ methanation by the use of AP-XPS.

3.4 Infrared spectroscopy

Infrared spectroscopy is a characterisation technique used to distinguish between different materials and to determine some of its constituents by measuring the IR absorption of the material. Unlike absorption of more energetic light, such as visible light, infrared light typically has a too low energy as to cause electronic transitions but instead suitably can excite vibrational states [41]. As such, IR light can be used to distinguish between different functional groups of a material based on their different absorption at different wavelengths. For absorption spectroscopy, it is foremost the transmission (T) and reflection (R) that is taken into regard whereas scattering effects are neglected. The quantitative expression of transmission spectroscopy follows equation (3.4) [41, 42]:

| Figure 3.3: A schematic figure showing the ejection of a photoelectron from the 1s orbital of oxygen by an X-ray photon. KE refers to the kinetic energy whereas BE is the binding energy. ϕ is the work function of the instrument. |
| Figure 3.4: Schematic representation of transmission and reflectance |
where the intensity of the incident light ($I_0$) is divided by the intensity of the transmitted ($I_T$) light. In the case of a reflection measurement the intensity of the reflected ($I_R$) light replaces $I_T$ in equation (3.4). The absorption of the light is expressed as absorbance ($A$), which is either the logarithm of the reciprocal transmission or reflectance of the light as expressed by equation (3.5) for transmission.

$$A = \log_{10} \frac{1}{T}$$

When measuring the absorbance, typically either transmission or reflectance is accounted for and in the case of a reflectance measurement, the absorbance is often referred to as a pseudo-absorbance.

**Diffuse reflectance**

There are two kinds of reflections, the specular and diffuse reflection out of which it is perhaps specular reflection that comes to one’s mind when thinking about reflection of light. Specular reflection is the mirror-like reflection. For specular reflection, the light is reflected at the same angle as the incident light in regards to the normal, giving a mirror-like or glossy appearance. This kind of reflection occurs for finely ordered materials such as larger single crystals but also flat and smooth materials such as metals and glasses. In contrast to this, diffuse reflection produces, as its name suggests, a more randomly distributed reflection. While the light is reflected at the same angle as the incident light (in regards to the norm) like the case of specular reflection, the incident rays do not share a common norm due to the polycrystalline structure and/or the roughness of the surface that creates a multitude of different reflection angles in regards to the average norm [42].

In contrast to transmission and specular reflection experiments, diffuse reflectance can easily be performed on a powder sample without any sample preparation as the phenomena that often plagues these two methods is instead the point of interest – the diffuse reflection.
Molecular vibrations and selection rules

While absorption of visible and ultraviolet light typically leads to electronic transitions, i.e., one or more electrons absorb the light and achieve a more energetic state, IR light can rarely produce electronic transitions but instead induce vibrations in the absorbing molecule. The energy for the promotion of this vibration is based on factors such as binding energies and bond angles, which can be realised by visualising the molecule as a harmonic oscillator [41, 43].

However, not all kinds of groups can be observed by IR spectroscopy due to selection rules. While it’s only possible for a material to absorb visible light if the energy of the photon corresponds to the energy gap between the electronic states, IR light has an extra criteria that origins from the molecular structure of the probed material. In order for a molecule to absorb light, the resulting vibration needs to induce a net change in the dipolar moment [43]. Take for example CO$_2$, there are two different possible stretchings, the symmetric and asymmetric stretching of the C=O bonds. While the symmetric stretch causes a change in the dipole moment for each isolated bond, there is no net change due to the two stretchings counteracting one another. Whereas for the asymmetric stretching, the two C=O vibrations change the dipole moment in the same direction, thus adding their contributions to a net change.

Table 3.1: The six normal modes of vibration, applied to the CO$_2$ molecule, with the net change in dipole moment, $\Sigma \vec{\mu}_i$. Note that of these 6 vibrations only 4 are observed for CO$_2$ as $\rho$ and $\tau$ are rotations for CO$_2$ and $\nu_s$ is IR inactive due to lack of net change in dipole moment.

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<th>Stretching</th>
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<td>$\nu_s$ (Symmetric)</td>
<td>$\delta$ (Scissoring)</td>
<td>$\omega$ (Wagging)</td>
</tr>
<tr>
<td>$\nu_{as}$ (Asymmetric)</td>
<td>$\rho$ (Rocking)</td>
<td>$\tau$ (Twisting)</td>
</tr>
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\[ \Sigma \vec{\mu}_i = 0 \]
\[ \Sigma \vec{\mu}_i > 0 \]
\[ \Sigma \vec{\mu}_i > 0 \]
\[ \Sigma \vec{\mu}_i = 0 \]
\[ \Sigma \vec{\mu}_i > 0 \]
\[ \Sigma \vec{\mu}_i = 0 \]

3.5 Modulation excitation spectroscopy

One of the greatest challenges when to determine the reaction pathway for a given reaction and catalyst is to distinguish true intermediates from spectators species that form during the reaction but not take part in the main reaction. One commonly utilised experimental design as to resolve this problem is the modulation
The figure shows a conceptual sketch of a MES experiment. Behaviour and content of the system which otherwise would be hidden can be determined indirectly by studying its response to a stimulating signal.

**Reaction steps of the simulated reaction**

\[
\begin{align*}
A + \cdot & \leftrightarrow A^* \quad (3.6) \\
B_2 + 2\cdot & \leftrightarrow 2B^* \quad (3.7) \\
A^* + B^* & \leftrightarrow C^* + \cdot \quad (3.8) \\
C^* + B^* & \leftrightarrow E^* + \cdot \quad (3.9) \\
E^* & \leftrightarrow E + \cdot \quad (3.10) \\
A^* + B^* & \leftrightarrow D^* + \cdot \quad (3.11) \\
A^* & \leftrightarrow A' + \cdot \quad (3.12)
\end{align*}
\]

* Empty adsorption site

\[
\theta_i = \frac{\text{Nr of sites occupied by } i}{\text{Total amount of sites}} \quad (3.13)
\]

As an example, consider the hypothetical reaction \(A + B_2 \longrightarrow E\) that is heterogeneously catalysed (solid catalyst, gaseous reactants). The intermediate reaction steps are given by equation (3.6) to (3.10).

In addition to these reactions, both a rapid and a slow side reaction occur as expressed by equation (3.11) and (3.12), respectively. In order to simplify the model, assumptions are made that include no presence of mass transport limitations, all the reactions occur on an extended flat surface and all adsorbate species except \(A'\) are adsorbed on the same type of adsorption site. By applying a modulation in the form of sinusoidal change of the partial pressure of \(A(p_A)\) on this reaction system while keeping the partial pressure of \(B\) constant, a response similar to that shown in Figure 3.8 can be obtained. While initially showing a response to the stimulus, the side reaction (3.12) eventually reaches steady-state due to its slow kinetics. This kind of response might be expected from stable, non-interactive species such as carbonates and hydroxides. While these can be easily filtered out by the modulation excitation spectroscopy (MES) approach, rapid reactions such as (3.11) may still show a response. A relevant example to this work would be the equilibrium reaction \(\text{HCOO} \rightleftharpoons \text{CO} + \text{OH}\).

**Phase-sensitive detection**

For data obtained from modulation excitation measurements, removal of the static signal and extraction of the delay for the responding signal compared to the modulation is a main target. In order to accomplish this, so-called phase sensitive detection (PSD) can be applied, either by Fourier transform or, as in this study, by a lock-in amplifier, expressed in regards to a transient IR data set in equation (3.14) [45]. In equation (3.14), the \(A(\tilde{\nu}, \tau)\) term is the amplitude in the time-domain, \(k\omega^1\) is the analysed frequency, \(\tau\) is
the period time, $\phi_k^{\text{PSD}}$ is the operator controlled phase angle and $A_0^{\text{PSD}}$ is the phase-resolved amplitudal data.

\[ A_k^{\phi_k^{\text{PSD}}}(\tilde{\theta}) = \frac{2}{\tau} \int_0^\tau A(\tilde{\theta}, t) \sin(k \omega t + \phi_k^{\text{PSD}}) dt \]  (3.14)

By application of equation (3.14) to MES data, signals that do not show a periodic response to the stimulus, such as noise and spectators, are reduced or fully cancelled out [46] and integration over additional periods further increases the signal-to-noise ratio. As an example, solving this on the previous simulated MES example (Figure 3.8) for the full phase period ($0^\circ \leq \phi_k^{\text{PSD}} \leq 360^\circ$) with the same frequency as the $p_A$ modulation results in Figure 3.9.

By finding the phase delay $\varphi_k$, i.e. how much out-of-phase the response is to the stimulus, one may sequence the order of events in a data set of multiple responses [47]. This phase delay can be obtained from the in-phase ($\phi_k^{\text{PSD}} = 0^\circ$) and out-of-phase ($\phi_k^{\text{PSD}} = 90^\circ$) amplitudes, as shown in (3.15) and (3.16) [45]. Solving the phase delays for the simulated example, one obtains $|\varphi_k|$ order of $D < A < C < E < B$, providing the correct reaction sequence ($A \rightarrow C \rightarrow E$).

\[ A_k(\tilde{\theta}) = \sqrt{A_k^0(\tilde{\theta})^2 + A_k^{90}(\tilde{\theta})^2} \]  (3.15)
\[ \cos \varphi_k = \frac{A_k^0(\tilde{\theta})}{A_k} , \quad \sin \varphi_k = \frac{A_k^{90}(\tilde{\theta})}{A_k} \]  (3.16)

As a square wave function may be expressed as a sum of infinite odd numbered harmonics (equation (3.17), Figure 3.10) one may study several frequencies from a single square wave MES experiment, given the corrections shown in equation (3.18) and (3.19) [47].

2: Note that the Nyquist-Shannon theorem still limits the highest studyable frequency to half the sampling frequency.
In this thesis MES-PSD has been employed to remove spectator signals from IR characterisation of adsorbates during in situ CO$_2$ methanation over Rh/CeO$_2$. Furthermore, by analysing phase delays, some proposals of reaction mechanisms are given.
Results and discussion

In regards to the study of the function of a catalyst, there are two interlinking aspects to consider — the behaviour of the catalyst material, and the surface reaction. In Paper I, we explored the change in crystal structure and oxidation state of the catalyst as well as formation of surface adsorbates. Due to some ambiguities regarding the adsorbates, a more in depth study focused solely on the surface reaction was performed in Paper II. In these studies, the catalyst was studied in the temperature span of 200-350 °C. The importance of this span, as reported in a previous kinetic study [48], is the initiation of CO (g) formation at temperatures between 250 and 350 °C whereas CH₄ (g) production occurs in the entire temperature interval.

4.1 Physico-chemical behaviour of the catalyst

In order to probe for the possibility of formation of new catalyst compounds during the reaction, such as oxides and carbides of the active metal, high-energy X-ray diffraction (HE-XRD) was performed. By this method the bulk phases of material domains with sufficient long range order can be studied in situ. Further, ambient pressure X-ray photoelectron spectroscopy (AP-XPS) was performed in order to observe changes in oxidation state of the surface in situ upon exposure to reaction atmospheres.

Observing the crystal structural dynamics

The crystal structural dynamics of the Rh/CeO₂ catalyst was studied by transient in situ HE-XRD using a modulation excitation approach (see Section 3.5). While the sample was exposed to a constant CO₂ flow (0.5 vol.%), the H₂ concentration was periodically changed between 0 and 2 vol.% in a square-wave fashion (20 min period time) while the diffractogram of the sample was recorded. The resulting diffractograms can be seen as difference plots in Figure 4.2. While weak, a peak corresponding to supported rhodium metal (Rhₚ) is observed during these experiments. This reflection corresponds to a d spacing of 2.2 Å, whereas 2.1 Å is expected for metallic Rh, which is likely due to a strong interaction with CeO₂ and possibly also with a boundary RhOₓ phase as observed with AP-XPS (below). As for the lack of strong rhodium

Figure 4.1: High-resolution TEM image of the Rh/CeO₂ catalyst used in the in situ structural study. The inset shows a 2 nm Rh crystallite, showing the 2.2 Å interplanar distance of the (111) plane.
peaks, this is likely due to a combination of amorphous rhodium metal and small particles (of which particle diameters of 2-3 nm were primarily observed in TEM and the average was 4 nm), for which both of these factors create a lack of well defined long range order. While no change in regards to rhodium is observed during this experiment, a decrease in $q$ for the reflections of CeO$_2$ during CO$_2$ hydrogenation could be clearly seen. This decrease in $q$ corresponds to an increase in interplanar distances ($d$), which in turn suggests an expansion of the unit cell. While this reduction is expected, the partial re-oxidation of CeO$_2$ following the removal of H$_2$ (increase in $q$) from the gas stream provides a new insight to the reaction. Namely, it suggests that either CO$_2$ itself or remaining reaction intermediate(s) from the hydrogenation reaction to some extent heals the oxygen vacancies formed at CeO$_2$. This shows that the redox behaviour of CeO$_2$ could be involved in the catalytic reaction mechanism, its role in the CO$_2$ methanation and rWGS reactions cannot be determined from this experiment alone.

### Changes in the surface oxidation state

The oxidation state of the catalyst surface was explored using AP-XPS. As seen in Figure 4.3, both RhO$_x$ and Rh metal phases with alternating ratios of the oxidised and reduced state are observed when the Rh/CeO$_2$ sample held at 595 K is exposed to a hydrogen (1.33 × 10$^{-6}$ hPa) and oxygen (1.33 × 10$^{-5}$ hPa) environment, respectively. Furthermore, the rhodium is not fully reduced under the hydrogen conditions. Similar to what is observed in the HE-XRD measurements, a partial reduction of CeO$_2$ is seen in the reducing environment as a decline of Ce$^{4+}$ signal.

As for in situ study of the methanation using AP-XPS, a transient study was performed in which the sample was exposed to a gas mixture of 0.0133 hPa CO$_2$ and 0.1867 hPa H$_2$ and the temperature was gradually increased from 515 to 625 K. Like the previous reduction-oxidation experiment, rhodium remains as partially oxidised during the experiment. As for the C 1s region, formation of surface species such as HCOO and CO is seen. While present on the surface at 515 K, no CO (g) was detected by the mass spectrometer while CH$_4$ production was seen during the whole temperature range. The first CO (g) could not be observed until 615 K was reached.

### 4.2 Exploring the surface reactions

While the previous AP-XPS results show that CO and HCOO species form on the Rh/CeO$_2$ catalyst, the possibility of them...
being intermediates could not be ascertained. As such, this was followed by a study focused solely on the surface reactions by the use of diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS). An initial assessment of the surface adsorbates during reaction conditions was performed by steady-state hydrogenation at 200, 250 and 300 °C. Afterwards, the hydrogenation was studied transiently at the same temperature in order to deduce which of the adsorbates that correspond to intermediates.

Active and inactive, adsorbates at steady-state

After exposing Rh/CeO$_2$ to a flow of 0.5 vol.% CO$_2$, carbonate (CO$_3$) formation is observed out of which bidentate (b-CO$_3$) is the primary mode of adsorption of CO$_2$ to CeO$_2$, see Figure 4.5a. After the introduction of 2 vol.% H$_2$ to the CO$_2$ stream, additional peaks with high absorption appeared. Based on that these peaks correspond to hydrogenation products and the previous observation of HCOO in the AP-XPS experiment, one would expect that the peaks in the 1600-1300 cm$^{-1}$ would correspond at least partially to HCOO. However, although C–H vibrations for HCOO are weaker than those of O–C–O, the weak C–H vibrations in the 2900-2800 cm$^{-1}$ region signifies that HCOO only constitutes a minor portion of the peaks in the 1600-1300 cm$^{-1}$ region. As for the prominent peak at 1744 cm$^{-1}$, its high wavenumber suggests a carbonyl such as a bridged CO which is further supported by its temperature dependence, coinciding with the rWGS. While showing some discrepancy (reference value: 1725 cm$^{-1}$ [49]) to previous assignments, this peak is likely primarily composed of CO located at the interface of Rh and CeO$_2$ ($\nu$Rh-CeO$_2$(C–O)). As for the mode of coordination, it has been suggested that the CO adopts a Rh–C=O–Ce coordination in which the oxygen is coordinated to a Lewis acid site of the support [49]. As for the higher wavenumber of this peak compared to the reference, this is possibly due to additional convoluted vibrations in the peak. Furthermore, minor peaks corresponding to species such as m-CO can be seen in the shoulder towards higher wavenumbers.

Figure 4.3: XPS spectra of Rh 3d (a) and Ce 4d (b) for reduced and oxidised Rh/CeO$_2$. Red squares in (b) is reduced Rh/CeO$_2$.

Figure 4.4: XPS spectra of Rh/CeO$_2$ during CO$_2$ methanation, showing C 1s during the temp. ramp (a) as well as C, Rh and Ce (b-d) at 515 K.
Figure 4.5: Steady-state in situ infrared spectra for the Rh/CeO$_2$ catalyst at the studied temperatures in the carbon-oxygen bond region 2200-800 cm$^{-1}$ during (a) CO$_2$ adsorption ($y_{CO_2} = 0.5$) and (b) hydrogenation ($y_{CO_2} = 0.5, y_{H_2} = 2$) as well as the complete measured region with a zoom at the C-H bond region 3000-2550 cm$^{-1}$ during hydrogenation (c).

Active species – to separate the wheat from the chaff

While providing an insight into the surface during reaction, it is difficult, if not impossible, to determine if an adsorbate is an intermediate or simply a spectator based on steady-state measurements. As such, in order to explore the possibility of these adsorbates being reaction intermediates, transient experiments with an modulation excitation approach were performed. Like the previous HE-XRD experiment, the CO$_2$ concentration in the gas flow was kept constant at 0.5 vol.%, whereas the hydrogen concentration was modulated in a square wave fashion, alternating between 0 and 2 vol.%. However, unlike the HE-XRD, a higher sampling frequency ($\sim$2 Hz) and shorter modulation periods ($\tau$) were employed in order to study the rapid reactions. Furthermore, both the modulation period ($\tau = 30, 60, 120$ s) as well as temperature ($T = 200, 250, 300$ °C) were varied. As an example, the case of 300 °C with a modulation period of 120 s can be seen in Figure 4.6 with the gas modulation (a), the infrared spectra (b) and the difference infrared spectra (c). As seen in Figure 4.6c, clear response from vibrations in the carbonyl (2200-1700 cm$^{-1}$), carbonate (1700-800 cm$^{-1}$) as well as hydroxide and C–H (3800-2700 cm$^{-1}$) region were obtained. In regards to the other experiments, while the same regions showed a response, different intensities (both absolute and relative between the peaks) were obtained, for which high temperature and long periods favoured the carbonyl region in particular. Furthermore, steady-state condition was reached for several peaks at 120 s during H$_2$ flow. However, the hydrogenation products were never depleted during the H$_2$ purge steps. This is at least partially due to the dead volume of the reaction cell which prohibits a full purge of hydrogen from the gas phase during the modulation period. As such, one should take into account that the reaction is only perturbed by the modulation, not initiated and quenched.

In order to more easily compare the adherence of the response
to the stimulus, remove static contributions as well as noise this
time-resolved data was further analysed by phase sensitive detec-
tion (PSD), using a lock-in amplifier (see equation (3.14)) which
was integrated over several periods. The obtained phase-resolved
spectra ($A^{\delta\text{PSD}}$) of the 2200-900 cm$^{-1}$ region can be observed in the
top rows of Figure 4.7 for modulation period 120 s (panel a), period
60 s (panel b), and period 30 s (panel c) between $0^\circ \leq \theta^{\text{PSD}} \leq 180^\circ$.
Furthermore, the bottoms rows shows a composite plot in which
the maximum amplitude ($A^{\delta\text{PSD}, \text{max}}$) is given at each wavenumber,
given a full period of $0^\circ \leq \theta^{\text{PSD}} \leq 360^\circ$, together with the corres-
ponding phase delay ($\varphi$) for the most pronounced peaks, marked
with bullets ($\bullet$).

Carbonyls

While minor peaks such as m-CO at 200 cm$^{-1}$ are seen, it is the
previously assigned $v_{\text{Rh-CeO}_2}$(C=O) vibration, from steady-state
 measurements, that is the most prominent peak in the MES-PSD
results, given $\tau \geq 60$ s and $T \geq 250$ °C. However, while appearing
as a single peak during steady-state, the MES-PSD results show
that it consists of at least two separate peaks approximately located
at 1700 and 1800 cm$^{-1}$. This is most noticeable in the $\varphi^{\text{PSD}, \text{max}}(\bar{\nu}, T =
200$ °C, $\tau = 60$ s) spectrum (Figure 4.7b). Furthermore, it is unlikely
that these vibrations belong to the same molecule as they do not
share a phase delay (i.e. they are not formed at the same time). Out
of these, the peak at 1700 cm$^{-1}$ is expected to correspond to the
previously assigned $v_{\text{Rh-CeO}_2}$(C=O) vibration. As for the 1800 cm$^{-1}$,
its low wavenumber makes it unlikely to correspond to a bridged
CO (br-CO) and is instead likely to correspond to CO in a hollow
or hollow-like site (h-CO)[50, 51]. While this convoluted peak is
Results and discussion

(a) Modulation period $\tau = 120$ s

(b) Modulation period $\tau = 60$ s

(c) Modulation period $\tau = 30$ s

Figure 4.7: Graphical representation of phase sensitive detection (PSD) analysis results for CO$_2$ hydrogenation over Rh/CeO$_2$ at 200, 250 and 300 °C and different H$_2$ modulation period lengths ($\tau$) of 120, 60 and 30 s. Top rows: PSD plots showing amplitude ($A^{\phi_{PSD}}$) versus wavenumber for demodulated spectra between $0^\circ \leq \phi_{PSD} \leq 180^\circ$. Bottom rows: Composite plots showing maximum amplitude ($A^{\phi_{PSD, \text{max}}}$) at each wavenumber in the period of $0^\circ \leq \phi_{PSD} \leq 360^\circ$ (--) together with the phase delay ($\phi$) of the peaks (●) and visual guides (---).
4.2 Exploring the surface reactions

primarily located close to the 1800 cm\(^{-1}\) peak, it shifts towards lower wavenumbers with increasing temperature. While a shift of the h-CO peak towards lower wavenumbers could be due to changes to its surrounding environment, perhaps a larger contributing factor is the increase of the CO\(_{\text{RhCeO}_2}\) to h-CO ratio. Furthermore, the intensity drop of these carbonyls, compared to other peaks in the spectra, at 300 °C coincides with the formation of CO (g) and bridged CO (br-CO, 1875 cm\(^{-1}\)). Although this decrease in h-CO could correspond to a potential increase in consumption rate of the hydrogenation reaction, it may also be due to an increase in desorption rate (giving a significant CO (g) formation) or a change in coordination to br-CO due to restructuring of the Rh surface. As for the relation between CO\(_{\text{RhCeO}_2}\) and h-CO, as CO\(_{\text{RhCeO}_2}\) the lower phase delay of CO\(_{\text{RhCeO}_2}\) opens up several different possibilities in regards reaction mechanisms. As an example, these carbonyls might be dependent on each other such as CO migration from the Rh-CeO\(_2\) site to the extended Rh surface. As such, while the Rh surface alone should be able to dissociate oxygen from CO\(_2\), the introduction of this new active site with faster kinetics could be linked to the promotional effect of CeO\(_2\) to Rh. However, one need to take into account that CO\(_{\text{RhCeO}_2}\) and h-CO do not share adsorption site and possibly not active site either. Factors such as ratio of adsorption sites that participate in the reaction contra only facilitate migration as well as competition with other species in regards to surface coverage will have an effect. As such, this needs to be further explored in order to support or disregard this idea.

<table>
<thead>
<tr>
<th>Species</th>
<th>(\tilde{\nu}) [cm(^{-1})]</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>(\nu(C=O))</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CO(_{\text{RhCeO}_2})</td>
<td>1700</td>
<td>[49]</td>
</tr>
<tr>
<td>h-CO</td>
<td>1800</td>
<td>[50]</td>
</tr>
<tr>
<td>br-CO</td>
<td>1875</td>
<td>[52]</td>
</tr>
<tr>
<td>m-CO</td>
<td>2010</td>
<td>[52]</td>
</tr>
<tr>
<td>(p)-CO(_3)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(\nu_{\text{as}}(\text{OCO}))</td>
<td>1460</td>
<td>[53]</td>
</tr>
<tr>
<td>(\nu_{\text{s}}(\text{OCO}))</td>
<td>1390</td>
<td>[53]</td>
</tr>
<tr>
<td>(b)-CO(_3)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(\nu_{\text{as}}(\text{OCO}))</td>
<td>1590</td>
<td>[54]</td>
</tr>
<tr>
<td>(\nu_{\text{s}}(\text{OCO}))</td>
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<td>[54]</td>
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<td>[55]</td>
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<tr>
<td>(\nu_{\text{s}}(\text{OCO}))</td>
<td>1300</td>
<td>[55]</td>
</tr>
<tr>
<td>(\nu(\text{CH}))</td>
<td>2850</td>
<td>[55]</td>
</tr>
<tr>
<td>(\delta(\text{CH}))</td>
<td>2950</td>
<td>[55]</td>
</tr>
</tbody>
</table>

### Carbonates and formates

While formates are present on the catalyst, based on the O–C–O vibrations at 1550 and 1300 cm\(^{-1}\) as well as C–H vibrations at 2950 and 2850 cm\(^{-1}\) seen in Figure 4.7 and 4.8 [55], they do not exhibit a strong response to the hydrogen stimulus. However, as very minor responses might simply be due to experimental fluctuations, such as disturbances to the total flow during a gas switch, it is possible that this response does not originate from a participation in the methanation reaction.

Carbonates, which often are too stable to participate in hydrogenation do show a response to the hydrogen stimulus. One such carbonate is the b-CO\(_3\) which was previously seen in the steady-state measurement. The b-CO\(_3\) showed a huge phase delay to the modulation (\(|\phi| > 180^\circ\)), signifying a consumption during hydrogen flow rather than production, as one would expect. However, another pair of vibrations are seen in the O–C–O region, those of 1460 and 1390 cm\(^{-1}\), instead show a minute phase delay. One would therefore expect these to correspond to hydrogenation products,
such as perhaps an inorganic carboxylate. However, the low $\nu_{\text{as-s}}$ split as well as the low wavenumber of the peaks makes this doubtful and they instead more likely correspond to polydentate carbonates. As the ceria surface reduces and restructures during the H$_2$ step, a p-CO$_3$ configuration might be more favourable for CO$_2$ adsorption rather than the previous b-CO$_3$ in the oxidised state. As such, the activity of these carbonates might therefore not be linked to the methanation reaction but are instead a response to the changes in the state of ceria. To gain an insight into the formation of p-CO$_3$, the phase delay of the carbonates was compared. By adjusting the phase delay of b-CO$_3$ with half a period ($\phi(b\text{-CO}_3) + 180 \approx -80^\circ$), it is revealed that p-CO$_3$ has a shorter phase delay (-30°) and its formation should therefore be an earlier event than the b-CO$_3$ consumption. As such, the primary mode of formation for p-CO$_3$ should therefore be CO$_2$ adsorption rather than re-configuration of b-CO$_3$.

### 4.3 Reaction mechanisms – formulating an idea

While a multitude of different reaction pathways for CO$_2$ methanation over supported metal catalysts could be theorised, only a few pathways have shown a likelihood. Of these, it is primarily the formate and carbon monoxide routes, named after their primary reaction intermediates, which has been observed experimentally.

During reaction conditions, while a portion of the rhodium is oxidised its surface is expected to be reduced whereas both the
bulk and surface of ceria adopts a partially reduced structure. A number of active species in the form of carboxyls, carbonates and formates could be seen. These carboxyls were active even at low temperatures where CO (g) formation has not been observed for Rh/CeO$_2$. As such, one might expect that the methanation on Rh/CeO$_2$ follows a carbon monoxide pathway. However, some activity was also observed from formate. While not excluding the carbon monoxide pathway, it does suggest that several pathways might be involved and perhaps interact with each other, by reactions such as HCOO $\rightleftharpoons$ CO + OH.

The Ce$^{4+}$/Ce$^{3+}$ cycling of CeO$_2$ could be a contributing factor in the promotion of CeO$_2$ for both methanation as well as rWGS. As was seen, CO formation at the Rh-CeO$_2$ interface as well as surface carbonates responded to the hydrogen stimulus. A potential mechanism could be the activation of CO$_2$ by an oxygen vacancy on the CeO$_2$ surface which is later on hydrogenated by H$_2$ spill-over from the Rh metal. A similar mechanism has been proposed for other catalysts[56], in which the oxygen vacancy is formed and healed in the reaction cycle. Similar behaviour was observed in the HE-XRD results, in which CeO$_2$ was partially re-oxidised during when H$_2$ was removed from the gas stream.

While leaving gaps for further exploration, the results of this work have provided an insight into the mechanisms behind the reaction pathway of CO$_2$ methanation over Rh/CeO$_2$. 
Conclusions and future work

The structural changes of the Rh/CeO$_2$ catalyst and its surface reactions during CO$_2$ methanation were studied \textit{in situ} by HE-XRD, AP-XPS and DRIFTS. It was revealed that CeO$_2$ takes an active role, cycling between Ce$^{4+}$ and Ce$^{3+}$. Presumably, this is due to the formation of oxygen vacancies which are further on partially healed by an intermediate involving reaction. Furthermore, while the rhodium was shown to be predominately in a reduced state during the reaction, a RhO$_x$ phase was observed. This RhO$_x$ could not be removed by reduction and is likely located at the intermediary region of Rh and CeO$_2$, caused by a strong metal support interaction.

The presence of active carbon monoxides on the catalyst strongly suggests that the catalyst follows a carbon monoxide pathway, possibly the carbide pathway. The Ce$^{4+}$ and Ce$^{3+}$ of CeO$_2$ affected the configuration of adsorbed carbonates, alternating between b-CO$_3$ and p-CO$_3$. If these carbonates also take part in the methanation reaction remains unknown. While some activity of formate could be observed, this is possibly due to an equilibrium reaction such as HCOO $\rightleftharpoons$ CO + OH rather than a support for the formate pathway being present. Furthermore, the formation of this formate could have been alleviated by CO$_2$ activation by the reduced CeO$_2$, possibly involving a short-lived bicarbonate (HCO$_3$) intermediate as reported for other catalysts [25].

5.1 Remaining questions

While a greater insight into the mechanisms of CO$_2$ methanation on Rh/CeO$_2$ has been gained, the complete reaction pathway remains unknown. Several kinds of surface carbon monoxide species were identified but their interactions, or lack thereof, to each other are not clear. As such, it cannot be stated whether or not only a few participate in a reaction step while the others respond due to processes such as reconfiguration and migration. Furthermore, the active sites remain elusive. The occurrence of CO in the rhodium-ceria interface suggests that CO$_2$ dissociation may occur in this catalytic site environment. If hydrogenation also occur on these sites or if the CO migrates to another site for hydrogenation is still unknown. A study of a potential migration on a model catalyst would therefore be of interest.

In this work, the MES-DRIFTS was performed using H$_2$ modulation in order to repeat the previous work in which the structural changes were characterised. While allowing one to find active species, the sequencing of the reaction pathway becomes daunting due to hydrogen being involved in several reaction steps. As
such, one should follow this study with a CO$_2$ modulated MES study as the linear path of C greatly simplifies the process.

Given that the pathway is determined, the determination of the limiting step(s) would be of interest. For instance, how does ceria promote the hydrogenation on rhodium? Is it primarily an electronic promotion or is it some other process, such as CO$_2$ dissociation at the Rh-CeO$_2$ interface, which provides the greatest beneficial effect? This is a highly complex question which is not easily answered by experiments alone.
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Felix Hemmingsson, Göteborg, May 2020
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Notation

$\delta(X)$ Symmetric in-plane bending, scissoring, of group X

$\nu_s(X)$ Symmetric stretching of group X

$\nu_{as}(X)$ Asymmetric stretching of group X

$\phi_{PSD}^{\psi_k}$ The operator controlled phase angle/phase angle for demodulation for the $k^{th}$ harmonic frequency

$\tilde{\nu}$ Wavenumber, reciprocal wavelength

$\varphi_k$ Phase delay in regards to the $k^{th}$ harmonic frequency

$A_{PSD}^{\psi_k}$ The phase-resolved amplitudal data for a given operator controlled angle and harmonic frequency

$d$ Interplanar distance in a crystal structure

$y_i$ Partial gas composition in regards to substance $i$. Expressed as molar or volumetric percent of the gas that is constituted by $i$

DRIFTS Diffuse reflectance infrared Fourier transform spectroscopy

MES Modulation excitation spectroscopy

PSD Phase sensitive detection